



Attorney Docket No.: 6116.200-US

cur

0340 11

PATENT

Box Seq

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of: Keith Anderson et al.

Serial No.: 09/757,788

Group Art Unit: 1614

Filed: January 10, 2001

Examiner: To be assigned

Confirmation No. 8259

For: Transepithelial Delivery Of GLP-1 Derivatives

CERTIFICATE OF MAILING UNDER 37 CFR 1.8(a)

Box Sequence
Commissioner for Patents
Washington, DC 20231

Sir:

I hereby certify that the attached correspondence comprising:

1. Petition and Fee For Extension of Time (in duplicate)
2. Response to Notice to Comply with Sequence Rules
3. Sequence Listing
4. Copy of Notice to Comply with Sequence Rules
5. Disk Containing Sequence Listing

is being deposited with the United States Postal Service as first class mail in an envelope addressed to:

Box Sequence
Commissioner for Patents
Washington, DC 20231

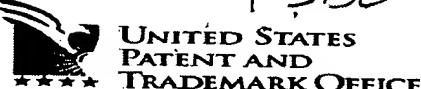
on December 11, 2002.

Rashida Haji
(name of person mailing paper)

Rashida Haji
(signature of person mailing paper)

RBOB/SBEN

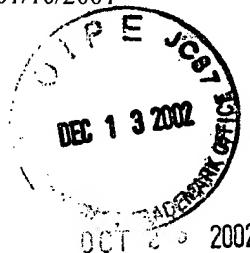
No 2211



Commissioner for Patents
Washington, DC 20231
www.uspto.gov

APPLICATION NUMBER	FILING/RECEIPT DATE	FIRST NAMED APPLICANT	ATTORNEY DOCKET NUMBER
09/757,788	01/10/2001	Keith Anderson	6116.200-US

Steve T. Zelson, Esq.
Novo Nordisk of North America, Inc.
Suite 6400
405 Lexington Avenue
New York, NY 10174-6400



CONFIRMATION NO. 8259
FORMALITIES LETTER



OC000000008941031

Date Mailed: 10/11/2002

**NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS
CONTAINING NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE
DISCLOSURES**

Filing Date Granted

This application contains sequence disclosures that are encompassed by the definitions for nucleotide and/or amino acid sequences set forth in 37 C.F.R. § 1.821(a)(1) and (a)(2). However, this application fails to comply with the requirements of 37 C.F.R. §§ 1.821-1.825. The application must be in sequence compliance before examination on the merits.

APPLICANT IS GIVEN ONE MONTH FROM THE DATE OF THIS LETTER WITHIN WHICH TO COMPLY WITH THE SEQUENCE RULES, 37 C.F.R. §§ 1.821-1.825. Failure to comply with these requirements will result in ABANDONMENT of the application under 37 C.F.R. § 1.821(g). Extension of time may be obtained by filing a petition accompanied by the extension fee under the provisions of 37 C.F.R. § 1.136. In no case may an applicant extend the period for response beyond the six-month statutory period. Direct the response to: The Office Of Initial Patent Examination.

See the attachment.

Applicant Must Provide as part of the response:

- An initial or substitute computer readable form (CRF) copy of the "Sequence Listing".
- An initial or substitute paper copy of the "Sequence Listing", as well as an amendment directing its entry into the specification.
- A statement that the content of the paper and computer readable copies are the same and, where applicable, include no new matter, as required by 37 C.F.R. 1.821(e) or 1.821(f) or 1.821(g) or 1.825(b) or 1.825(d).

For questions regarding compliance to these requirements, please contact:

- For Rules Interpretation, call (703) 308-4216
- To Purchase PatentIn Software, call (703) 306-2600
- For PatentIn Software Program Help, call (703) 306-4119 or e-mail at patin21help@uspto.gov or patin3help@uspto.gov

7b
A copy of this notice MUST be returned with the reply.

Customer Service Center

Initial Patent Examination Division (703) 308-1202

PART 2 - COPY TO BE RETURNED WITH RESPONSE